Study Number: IMM20704

Test Type: TOX Route: Oral Gavage

Species/Strain: Mouse/B6C3F1/N

C Number:

Study Gender:

PWG Approval Date

M12: Cytotoxic T Cell Activity **Test Compound:** Resveratrol

CAS Number: 501-36-0

Date Report Requested: 03/20/2020 Time Report Requested: 12:54:40

Lab: NTP

IMM20704

Male

See web page for date of PWG Approval

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Males: Study 2

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Treatment Groups (mg/kg/day)						
0	156	312	625	1250	2500	50 mg/kg CPS
15.56 ± 1.58 (8)	24.20 ± 3.31 (8)	27.50 ± 3.92 (8)	23.16 ± 5.71 (8)	17.48 ± 2.26 (8)	25.47 ± 4.07 (7)	-0.41 ± 0.22 (7) **
30.23 ± 2.86 (8)	42.12 ± 4.38 (8)	46.25 ± 4.70 (8)	38.52 ± 7.25 (8)	33.21 ± 3.27 (8)	44.95 ± 5.56 (7)	0.31 ± 0.93 (7) **
$49.76 \pm 4.70 (8)$	59.09 ± 4.54 (8)	63.62 ± 4.57 (8)	55.02 ± 7.53 (8)	50.42 ± 4.46 (8)	64.32 ± 6.11 (7)	-0.02 ± 0.37 (7) **
64.92 ± 4.37 (8)	74.57 ± 3.06 (8)	75.10 ± 3.98 (8)	66.79 ± 5.45 (8)	67.87 ± 4.69 (8)	78.37 ± 5.92 (7)	1.05 ± 0.82 (7) **
76.87 ± 2.44 (8)	86.09 ± 3.08 (8)	85.72 ± 3.67 (8)	82.03 ± 5.10 (8)	82.26 ± 3.44 (8)	89.11 ± 4.54 (7)	2.49 ± 1.38 (7) **
88.86 ± 2.14 (8)	93.80 ± 2.63 (8)	94.25 ± 3.25 (8)	91.54 ± 3.89 (8)	87.32 ± 3.64 (8)	95.55 ± 2.30 (7)	5.53 ± 2.62 (7) **
	$15.56 \pm 1.58 (8)$ $30.23 \pm 2.86 (8)$ $49.76 \pm 4.70 (8)$ $64.92 \pm 4.37 (8)$ $76.87 \pm 2.44 (8)$	$15.56 \pm 1.58 (8)$ $24.20 \pm 3.31 (8)$ $30.23 \pm 2.86 (8)$ $42.12 \pm 4.38 (8)$ $49.76 \pm 4.70 (8)$ $59.09 \pm 4.54 (8)$ $64.92 \pm 4.37 (8)$ $74.57 \pm 3.06 (8)$ $76.87 \pm 2.44 (8)$ $86.09 \pm 3.08 (8)$	0156312 $15.56 \pm 1.58 (8)$ $24.20 \pm 3.31 (8)$ $27.50 \pm 3.92 (8)$ $30.23 \pm 2.86 (8)$ $42.12 \pm 4.38 (8)$ $46.25 \pm 4.70 (8)$ $49.76 \pm 4.70 (8)$ $59.09 \pm 4.54 (8)$ $63.62 \pm 4.57 (8)$ $64.92 \pm 4.37 (8)$ $74.57 \pm 3.06 (8)$ $75.10 \pm 3.98 (8)$ $76.87 \pm 2.44 (8)$ $86.09 \pm 3.08 (8)$ $85.72 \pm 3.67 (8)$	Treatment Groups (mg/kg 0 156 312 625 15.56 ± 1.58 (8) 24.20 ± 3.31 (8) 27.50 ± 3.92 (8) 23.16 ± 5.71 (8) 30.23 ± 2.86 (8) 42.12 ± 4.38 (8) 46.25 ± 4.70 (8) 38.52 ± 7.25 (8) 49.76 ± 4.70 (8) 59.09 ± 4.54 (8) 63.62 ± 4.57 (8) 55.02 ± 7.53 (8) 64.92 ± 4.37 (8) 74.57 ± 3.06 (8) 75.10 ± 3.98 (8) 66.79 ± 5.45 (8) 76.87 ± 2.44 (8) 86.09 ± 3.08 (8) 85.72 ± 3.67 (8) 82.03 ± 5.10 (8)	Treatment Groups (mg/kg/day) 0 156 312 625 1250 15.56 ± 1.58 (8) 24.20 ± 3.31 (8) 27.50 ± 3.92 (8) 23.16 ± 5.71 (8) 17.48 ± 2.26 (8) 30.23 ± 2.86 (8) 42.12 ± 4.38 (8) 46.25 ± 4.70 (8) 38.52 ± 7.25 (8) 33.21 ± 3.27 (8) 49.76 ± 4.70 (8) 59.09 ± 4.54 (8) 63.62 ± 4.57 (8) 55.02 ± 7.53 (8) 50.42 ± 4.46 (8) 64.92 ± 4.37 (8) 74.57 ± 3.06 (8) 75.10 ± 3.98 (8) 66.79 ± 5.45 (8) 67.87 ± 4.69 (8) 76.87 ± 2.44 (8) 86.09 ± 3.08 (8) 85.72 ± 3.67 (8) 82.03 ± 5.10 (8) 82.26 ± 3.44 (8)	Treatment Groups (mg/kg/day) 0 156 312 625 1250 2500 15.56 ± 1.58 (8) 24.20 ± 3.31 (8) 27.50 ± 3.92 (8) 23.16 ± 5.71 (8) 17.48 ± 2.26 (8) 25.47 ± 4.07 (7) 30.23 ± 2.86 (8) 42.12 ± 4.38 (8) 46.25 ± 4.70 (8) 38.52 ± 7.25 (8) 33.21 ± 3.27 (8) 44.95 ± 5.56 (7) 49.76 ± 4.70 (8) 59.09 ± 4.54 (8) 63.62 ± 4.57 (8) 55.02 ± 7.53 (8) 50.42 ± 4.46 (8) 64.32 ± 6.11 (7) 64.92 ± 4.37 (8) 74.57 ± 3.06 (8) 75.10 ± 3.98 (8) 66.79 ± 5.45 (8) 67.87 ± 4.69 (8) 78.37 ± 5.92 (7) 76.87 ± 2.44 (8) 86.09 ± 3.08 (8) 85.72 ± 3.67 (8) 82.03 ± 5.10 (8) 82.26 ± 3.44 (8) 89.11 ± 4.54 (7)

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LEGEND

Data are displayed as mean ± SEM (N) unless otherwise noted.

Data displayed as a mean of (effector cell:target cell ratio)

CTL - Cytotoxic T Lymphocytes

^aCTL Activity is expressed as % target cell killing calculated as (sample Cr51 release - spontaneous Cr51 release / total Cr51 release - spontaneous Cr51 release)

Statistical analysis performed by Jonckheere (trend) and Shirley or Dunn (pairwise) tests.

Statistical analysis for the positive control group compared to the vehicle control group was performed using the Kruskal-Wallis test.

Statistical significance for the control group indicates a significant trend test

Statistical significance for a treatment group indicates a significant pairwise test compared to the vehicle control group

* Statistically significant at P <= 0.05

** Statistically significant at P <= 0.01

CPS = Cyclophosphamide

** END OF REPORT **